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Scientific and Technical Information Center

SEARCH REQUEST FORM

10/674,516

Requester's Full Name: Jeffrey E. Russel Examiner #: 62785 Date: 4-7-2005

Art Unit: 1654 Phone Number: 2-0969 Serial Number: ~~60/674,516~~

Location (Bldg/Room#): REM 3019 (Mailbox #): 3C18 Results Format Preferred (circle): (PAPER) DISK

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To ensure an efficient and quality search, please attach a copy of the cover sheet, claims, and abstract or fill out the following:

Title of Invention: Novel Peptide With Osteogenic Activity

Inventors (please provide full names): S. Ohnaraj, A. Gosiewska, A. Rezania, G. Heavner, X. Lin, C. Yi

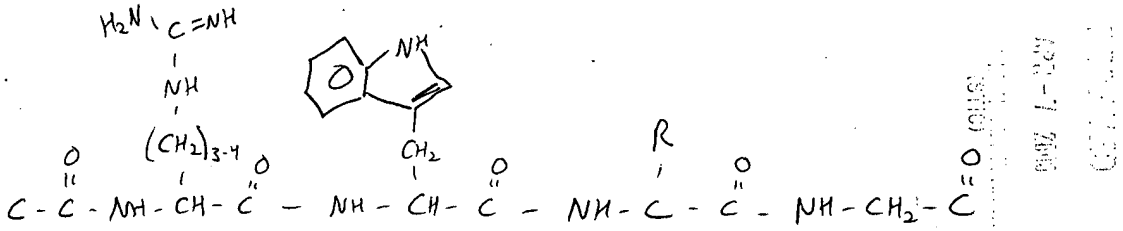
Earliest Priority Date: 9-10-2004

Search Topic:

Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc., if known.

For Sequence Searches Only ***Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.***

Please search the following partial structure:



where R is $-CH_2-SH$, $-CH_2-CH_2-SH$, or $-C(CH_3)_2-SH$

Thank you.

JER

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STAFF USE ONLY

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Searcher Phone #: _____

Searcher Location: _____

Date Searcher Picked Up: _____

Date Completed: _____

Searcher Prep & Review Time: _____

Online Time: _____

Type of Search

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_____ AA Sequence (#)

_____ Structure (#)

_____ Bibliographic

_____ Litigation

_____ Fulltext

_____ Other

Vendors and cost where applicable

_____ STN _____ Dialog

_____ Questel/Orbit _____ Lexis/Nexis

_____ Westlaw _____ WWW/Internet

____ In-house sequence systems

____ Commercial ____ Oligomer ____ Score/Length

____ Interference ____ SPDI ____ Encode/Transl

_____ Other (specify)

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L3             28 SEA SSS FUL L1

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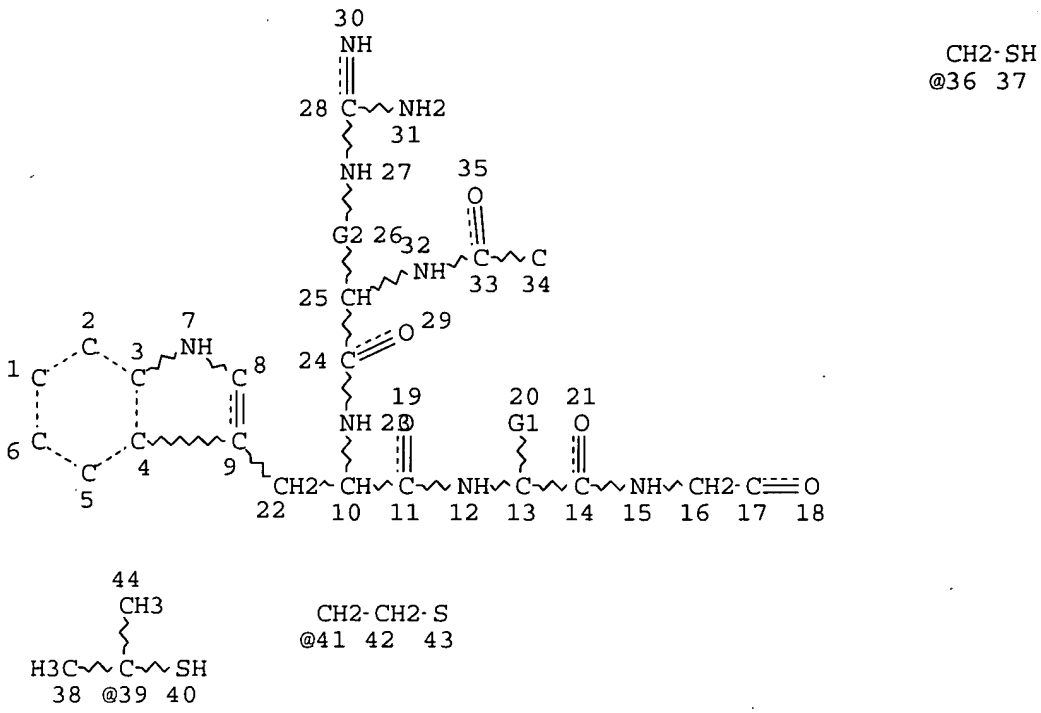


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L14 ANSWER 1 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:1060669 HCAPLUS
DOCUMENT NUMBER: 142:34829
TITLE: Knockout identification of target-specific sites in peptides by serial substitution of conformationally restricted metal-complexed residues in metallopeptide analogs
INVENTOR(S): Sharma, Shubh D.; Shi, Yi-Qun; Bastos, Margarita; Rajpurohit, Ramesh; Cai, Hui-Zhi
PATENT ASSIGNEE(S): Palatin Technologies, Inc., USA
SOURCE: U.S. Pat. Appl. Publ., 43 pp., Cont.-in-part of U.S. Ser. No. 464,117.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 4
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004248212	A1	20041209	US 2004-769695	20040130
WO 2002064734	A2	20020822	WO 2001-US50075	20011219
WO 2002064734	A3	20031120		

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US 2005014193	A1	20050120	US 2003-464117	20030617
WO 2004075830	A2	20040910	WO 2004-US2933	20040202

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PRIORITY APPLN. INFO.:		
US 2000-256842P	P	20001219
US 2001-304835P	P	20010711
US 2001-327835P	P	20011004
WO 2001-US50075	A1	20011219
US 2003-444129P	P	20030131
US 2003-464117	A2	20030617
US 2004-769695	A	20040130

AB The invention provides methods for identification and determination of target-specific sites in peptides and proteins, including a method for determining the primary sequence of a secondary structure within a known parent

polypeptide that binds to the target of interest. In one embodiment of the invention, a residue or mimetic containing a nitrogen atom and a sulfur atom available for binding to a metal ion is serially substituted for single residues in or inserted between adjacent residues in a known primary sequence of a peptide or protein. A residue or mimetic containing a nitrogen atom and a sulfur atom available for binding to a metal ion is serially substituted for single residues in or inserted between adjacent residues in a known primary sequence of the peptide or protein. The resulting sequence is complexed with a metal ion thereby forming a metallopeptide with a conformationally fixed and predictable secondary structure of the residues involved in metal ion complexation. The resulting metallopeptides are then used in binding or functional assays related to the target of interest, and the metallopeptide(s) which result in significant or substantially decreased or changed binding or functionality are determined to identify the primary sequence involved in such binding or functionality. The method is exemplified by α-MSH and bombesin analogs containing L-/D-cysteine insertions or substitutions complexed to the rhenium metal ion, and their binding to their resp. receptors.

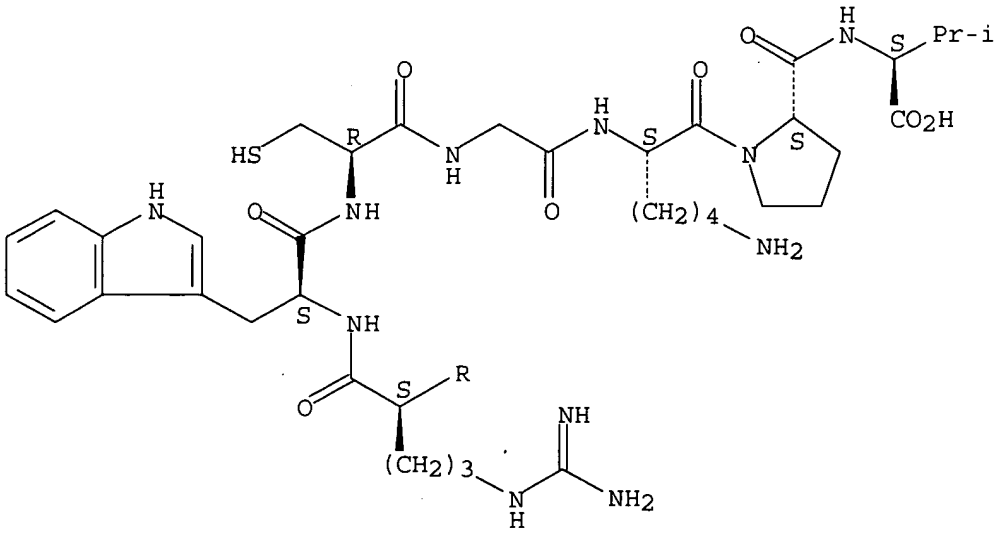
IT **754240-34-1D**, complex with rhenium metal ion **754240-92-1D**
, complex with rhenium metal ion
RL: ARU (Analytical role, unclassified); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study)
(knockout identification of target-specific sites in peptides by serial substitution of conformationally restricted metal-complexed residues in metallopeptide analogs)

RN 754240-34-1 HCAPLUS

CN L-Valine, L-seryl-L-tyrosyl-L-seryl-L-norleucyl-L-α-glutamyl-L-histidyl-L-phenylalanyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl-L-lysyl-L-prolyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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L14 ANSWER 2 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:740117 HCAPLUS
DOCUMENT NUMBER: 141:256945
TITLE: Knockout identification of target-specific sites in peptides by serial substitution of conformationally restricted metal-complexed residues in metallopeptide analogs
INVENTOR(S): Sharma, Shubh D.; Shi, Yi-Qun; Rajpurohit, Ramesh; Bastos, Margarita; Cai, Hui-Zhi
PATENT ASSIGNEE(S): Palatin Technologies, Inc., USA
SOURCE: PCT Int. Appl., 78 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 4
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
WO 2004075830	A2	20040910	WO 2004-US2933		20040202
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RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG					
US 2004248212	A1	20041209	US 2004-769695		20040130
PRIORITY APPLN. INFO.:			US 2003-444129P	P	20030131
			US 2004-769695	A	20040130
			US 2000-256842P	P	20001219
			US 2001-304835P	P	20010711
			US 2001-327835P	P	20011004
			WO 2001-US50075	A1	20011219

US 2003-464117 A2 20030617

AB The invention provides methods for identification and determination of target-specific sites in peptides and proteins, including a method for determining the primary sequence of a secondary structure within a known parent polypeptide that binds to the target of interest. A residue or mimetic containing a nitrogen atom and a sulfur atom available for binding to a metal ion is serially substituted for single residues in or inserted between adjacent residues in a known primary sequence of the peptide or protein. The resulting sequence is complexed with a metal ion thereby forming a metallopeptide with a conformationally fixed and predictable secondary structure of the residues involved in metal ion complexation. The resulting metallopeptides are then used in binding or functional assays related to the target of interest, and the metallopeptide(s) which result in significant or substantially decreased or changed binding or functionality are determined to identify the primary sequence involved in such binding or functionality. The method is exemplified by α-MSH and bombesin analogs containing L-/D-cysteine insertions or substitutions complexed to the rhenium metal ion, and their binding to their resp. receptors.

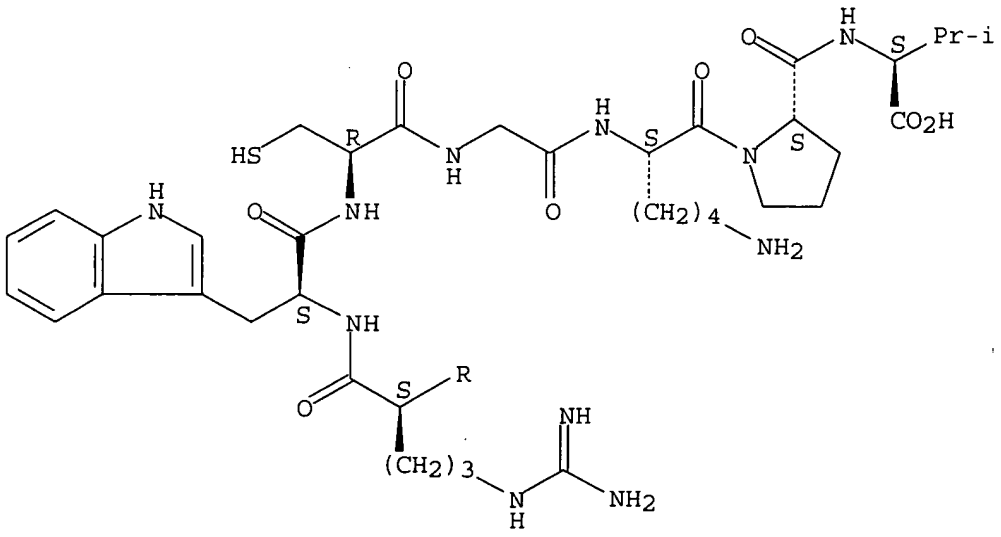
IT **754240-34-1D**, complex with rhenium metal ion **754240-92-1D**
, complex with rhenium metal ion
RL: ARU (Analytical role, unclassified); BSU (Biological study, unclassified); ANST (Analytical study); BIOL (Biological study)
(knockout identification of target-specific sites in peptides by serial substitution of conformationally restricted metal-complexed residues in metallopeptide analogs)

RN 754240-34-1 HCAPLUS

CN L-Valine, L-seryl-L-tyrosyl-L-seryl-L-norleucyl-L-α-glutamyl-L-histidyl-L-phenylalanyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl-L-lysyl-L-prolyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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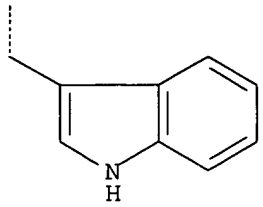
L14 ANSWER 3 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:310865 HCAPLUS
DOCUMENT NUMBER: 140:332438
TITLE: Method of screening for peptides capable of specifically acting on biological membrane and the pharmaceutical activities of the peptides
INVENTOR(S): Machida, Sachiko; Tokuyasu, Ken; Matsunaga, Shigeru; Sakakibara, Yoshikiyo; Kobori, Masuko; Wen, Zhesheng
PATENT ASSIGNEE(S): Japan
SOURCE: U.S. Pat. Appl. Publ., 58 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.		DATE
US 2004072992	A1	20040415	US 2003-651563		20030829
JP 2004248666	A2	20040909	JP 2003-303747		20030827
PRIORITY APPLN. INFO.:			JP 2002-253169	A	20020830
			JP 2003-21198	A	20030129

AB Peptide are provided, consisting of amino acid sequences comprising an amino acid sequence Z1X1X2X3Z2X4X5Z3X6X7X8Z4X9, where X1 -X9 are any amino acids and at least two amino acids of Z1-Z4 are basic amino acids, such that the peptides specifically act on biol. membranes of microorganisms and does not act on normal animal cell membranes. The peptides are selected from the group consisting of ALR, WALR, WGALR, RLAWG, GWALR RVL, KVL, RVG, KVG, GVR, VGR, RVA, RSV, RVS, KVS, SVK, and VSK, or an analog thereof. Screening for nucleic acids encoding a peptide capable of acting on a biol. membrane is achieved by constructing a DNA library, preparing peptides by cell-free transcription/translation, and selecting the peptides capable of specifically binding to a lipid bilayer membrane model immobilized on a solid phase such as a magnetic bead. The peptides demonstrate biol. activities in preventing putrefaction of food or industrial products, treating infectious disease, treating cancer, suppressing apoptosis, and killing microorganisms pathogenic to animals or plants (e.g., Erwinia carotovora).
IT **678998-11-3**

PAGE 2-A



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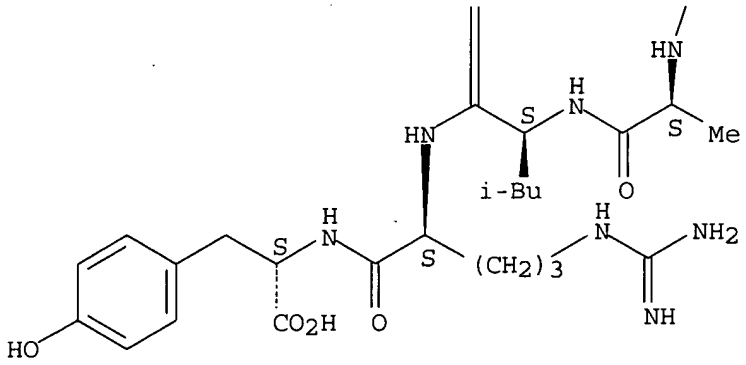

L14 ANSWER 4 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:20444 HCAPLUS
DOCUMENT NUMBER: 140:110119
TITLE: Mammalian EPO mimetic CH1 deleted mimetibodies, compositions, methods and uses for diagnosis and therapy of human diseases
INVENTOR(S): Heavner, George A.; Knight, David M.; Ghrayeb, John; Scallon, Bernard J.; Nesspor, Thomas C.; Kutoloski, Karen A.
PATENT ASSIGNEE(S): Centocor, Inc., USA
SOURCE: PCT Int. Appl., 123 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004002424	A2	20040108	WO 2003-US20495	20030630

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RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,

BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRIORITY APPLN. INFO.: US 2002-392431P P 20020628
US 2002-412144P P 20020919

AB The present invention relates to at least one novel erythropoietin (EPO) human CH1-deleted mimetibody or specified portion or variant, including isolated nucleic acids that encode at least one CH1-deleted mimetibody or specified portion or variant, CH1-deleted mimetibody or specified portion or variants, vectors, host cells, transgenic animals or plants, and methods of making and using thereof, including therapeutic compns., methods and devices. Peptides that mimic the activity of EPO, TPO, growth hormones, G-CSF, GM-CSF, IL-1, leptin, CTLA4, TRAIL, TGF-α, and TGF-β are the focus of this genetic engineering. The CH1 deleted mimetibody can optionally comprise at least one CH3 constant region directly linked with at least one CH2 region directly linked with at least one hinge region or fragment thereof directly linked with at least one partial V region, directly linked with an optional linker sequence, directly linked to at least one therapeutic peptide, optionally further directly linked with at least a portion of at least one variable antibody sequence. In a preferred embodiment a pair of a CH3-CH2-hinge-partial J sequence-linker-therapeutic peptide with an option N-terminal antibody sequence, the pair optionally linked by association or covalent linkage, such as, but not limited to, a Cys-Cys disulfide bond. The aim of the invention is use of the purified recombinant proteins for diagnosis or treatment of anemia, immune or autoimmune disease, cancer, or infectious diseases.

IT **268228-13-3**
RL: DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(amino acid sequence, mimetibody comprising; mammalian EPO mimetic CH1 deleted mimetibodies, compns., methods and uses for diagnosis and therapy of human diseases)

RN 268228-13-3 HCAPLUS

CN L-Serine, L-seryl-L-cysteinyl-L-tyrosyl-L-α-glutamyl-L-tryptophylglycyl-L-lysyl-L-leucyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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L14 ANSWER 5 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:20438 HCAPLUS
DOCUMENT NUMBER: 140:110118
TITLE: Mammalian CH1 deleted mimetibodies, compositions, methods and uses for diagnosis and therapy of human diseases
INVENTOR(S): Heavner, George A.; Knight, David M.; Ghrayeb, John; Scallon, Bernard J.; Nesspor, Thomas C.; Kutoloski, Karen A.
PATENT ASSIGNEE(S): Centocor, Inc., USA
SOURCE: PCT Int. Appl., 129 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004002417	A2	20040108	WO 2003-US20347	20030627
WO 2004002417	A3	20041104		

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PRIORITY APPLN. INFO.: US 2002-392431P P 20020628

AB The present invention relates to at least one novel human CH1-deleted mimetibody or specified portion or variant, including isolated nucleic acids that encode at least one CH1-deleted mimetibody or specified portion or variant, CH1-deleted mimetibody or specified portion or variants, vectors, host cells, transgenic animals or plants, and methods of making and using thereof, including therapeutic compns., methods and devices. In one embodiment, a CH1 deleted mimetibody comprises formula: (V1(n)-Pep(n)-Flex(n)-V2(n)-pHinge(n)-CH2(n)-CH3(n))(m), where V1 is at least one portion of an N-terminus of an Ig variable region, Pep is at least one bioactive peptide, Flex is polypeptide that provides structural flexibility by allowing the mimetibody to have alternative orientations and binding properties, V2 is at least one portion of a C-terminus of an Ig variable region, pHinge is at least a portion of an Ig variable hinge region, CH2 is at least a portion of an Ig CH2 constant region, CH3 is at least a portion of an Ig CH3 constant region, n and m can be an integer between 1 and 10. Peptides that mimic the activity of EPO, TPO, growth hormones, G-CSF, GM-CSF, IL-1ra, leptin, CTLA4, TRAIL, TGF-α and TGF-β are the focus of this genetic engineering. The aim of the invention is use of the purified recombinant proteins for diagnosis or treatment of anemia, immune or autoimmune disease, cancer, or infectious

diseases.

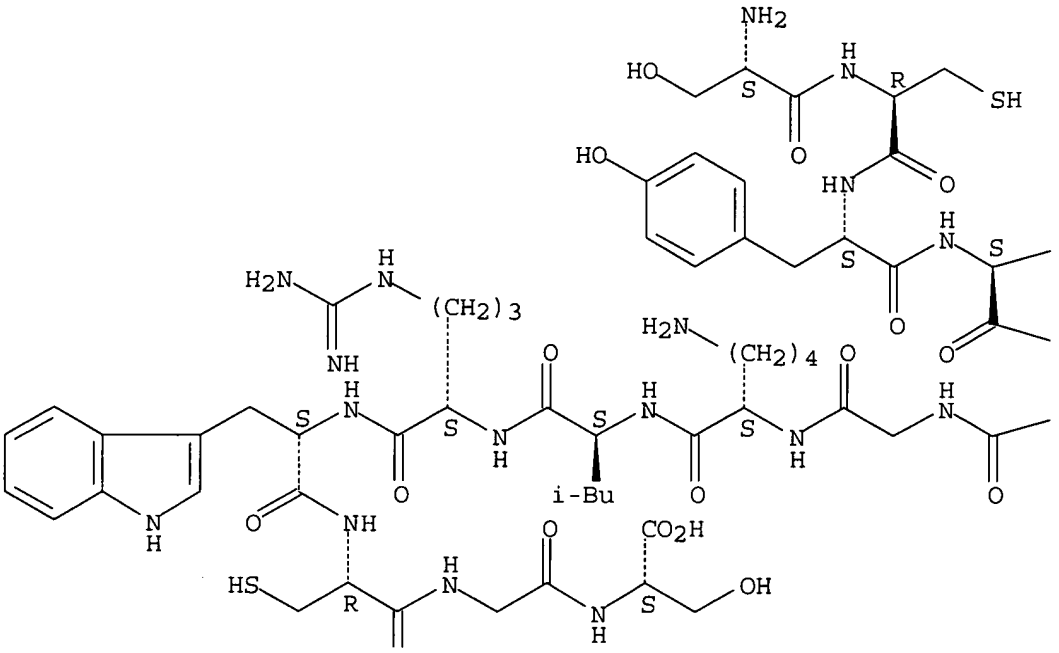
IT **268228-13-3**
RL: DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(amino acid sequence, mimetibody comprising; mammalian CH1 deleted mimetibodies, compns., methods and uses for diagnosis and therapy of human diseases)

RN 268228-13-3 HCAPLUS

CN L-Serine, L-seryl-L-cysteinyl-L-tyrosyl-L-α-glutamyl-L-tryptophylglycyl-L-lysyl-L-leucyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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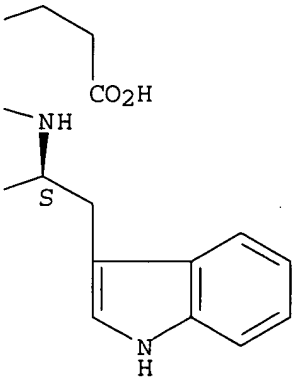


PAGE 2-A

L14 ANSWER 6 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2003:830375 HCAPLUS
DOCUMENT NUMBER: 139:303032
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human tissues
INVENTOR(S): Penn, Sharron Gaynor; Rank, David Russell; Hanzel, David Kagen
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 80 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 9
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2003194704	A1	20031016	US 2002-29386	20020403
US 2003194704	A1	20031016	US 2002-29386	20020403
PRIORITY APPLN. INFO.:			US 2002-29386 A	20020403

AB Methods and apparatus for predicting, confirming and displaying functional regions from genomic sequence data are used to identify unique human genome-derived single exon probes useful for gene expression anal., particularly gene expression anal. by microarray. Human BAC sequences

totaling .apprx.350 Mb of sequence (.apprx.10% of the human genome) were analyzed for exons using four sep. gene finding programs (GRAIL uses a neural network, GENEFINDER uses a hidden Markoff model, DICTION operates according to a different heuristic, and GENSCAN) and Mouse comparative genomics as a fifth gene prediction method. The exons were PCR amplified from genomic DNA, verified on agarose gels, and sequenced using universal primers to validate the identity of the amplicon to be spotted in microarrays. Thus, expression, homol., and functional information are provided for 13,700 unique genome-derived single exon probes are expressed at significant levels in one or more of 8 tested tissues: human brain, heart, fetal and adult liver, placenta, lung, bone marrow, and HeLa cells. The probes lack prokaryotic and bacteriophage vector sequences, as well as lacking homopolymeric stretches of A or T. Also presented are genome-derived single exon microarrays that include such probes, peptides encoded by the exons, and antibodies thereto. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of nine records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

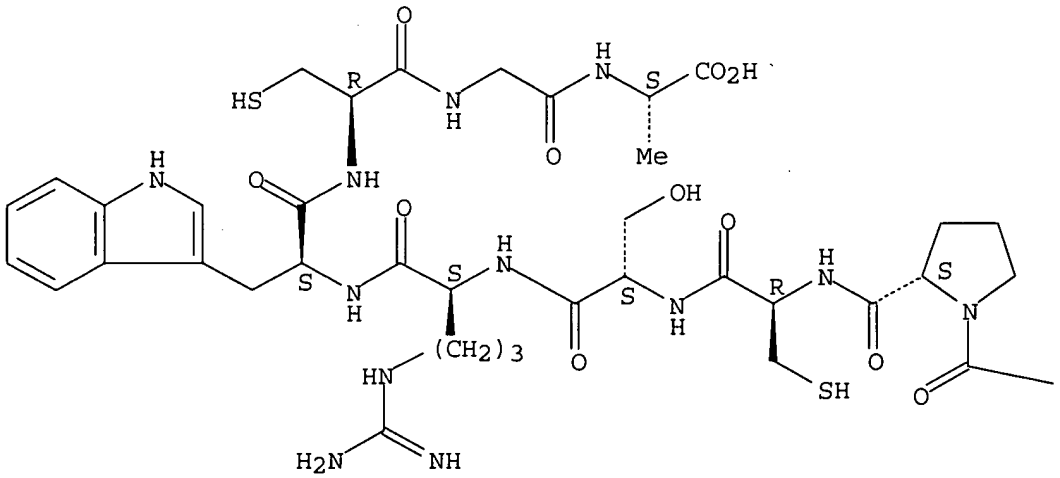
IT **610771-20-5 610774-96-4**
RL: ARG (Analytical reagent use); BSU (Biological study, unclassified); BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical study); BIOL (Biological study); USES (Uses)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human tissues)

RN 610771-20-5 HCAPLUS

CN L-Alanine, L-leucyl-L-arginylglycyl-L-prolyl-L-glutaminyl-L-seryl-L-prolyl-L-arginyl-L-cysteinyl-L-lysyl-L-prolyl-L-valyl-L-leucyl-L-valyl-L-histidylglycyl-L-glutaminyl-L-leucylglycyl-L-threonyl-L-arginyl-L-prolyl-L-cysteinyl-L-seryl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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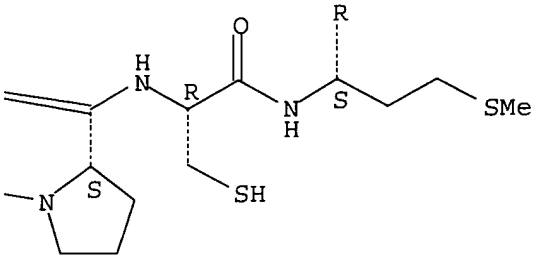

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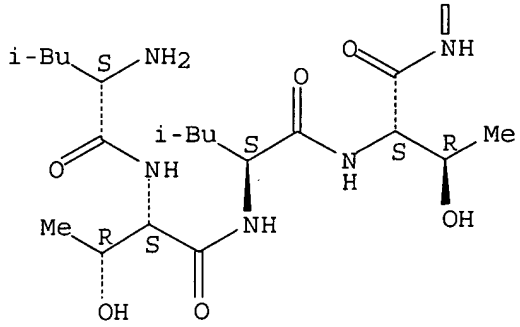
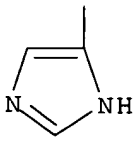
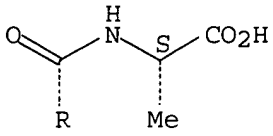






L14 ANSWER 7 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2003:818235 HCAPLUS
DOCUMENT NUMBER: 139:322283
TITLE: Methods for production and use of mammalian complementarity determining region mimetibodies for diagnosis and therapy of human diseases
INVENTOR(S): Heavner, George A.; Knight, David M.; Scallon, Bernard J.; Ghrayeb, John
PATENT ASSIGNEE(S): Centocor, Inc., USA
SOURCE: PCT Int. Appl., 97 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003084477	A2	20031016	WO 2003-US9139	20030324

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,

LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM,
PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT,
TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR,
BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2002-368791P P 20020329

AB This invention pertains to methods for production and use of mammalian complementarity determining region (CDR) mimetibodies for diagnosis and therapy of human diseases. Genetic engineering, expression, and purification of human mimetibodies containing Ig fragments (CDR, variable, framework and/or constant region) as well as a ligand binding domain are disclosed in this invention. Peptides that mimic the activity of EPO, TPO, growth hormones, G-CSF, GM-CSF, IL-1ra, leptin, CTLA4, TRAIL, TGF-α and TGF-β are the focus of this genetic engineering. The aim of the invention is use of the purified recombinant proteins for diagnosis or treatment of anemia, immune or autoimmune disease, cancer, or infectious diseases. At the time of publication, claimed sequence nos. 997 to 1109 were missing, and claimed sequence nos. 984 to 996 were not clearly identified.

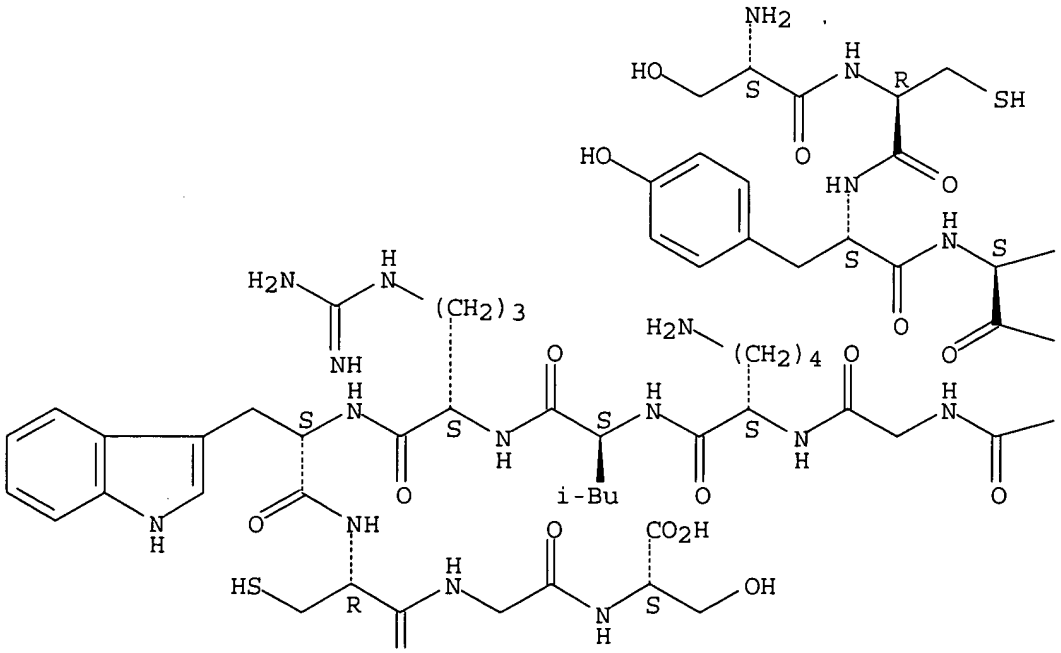
IT **268228-13-3**
RL: DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(calmodulin antagonist peptide; methods for production and use of mammalian CDR mimetibodies for diagnosis and therapy of human diseases)

RN 268228-13-3 HCAPLUS

CN L-Serine, L-seryl-L-cysteinyl-L-tyrosyl-L-α-glutamyl-L-tryptophylglycyl-L-lysyl-L-leucyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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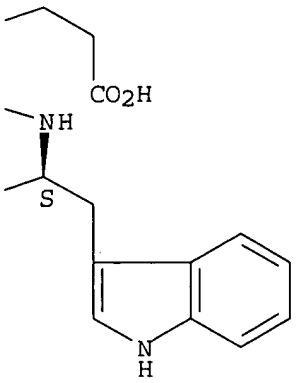


PAGE 2-A

L14 ANSWER 8 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2003:320039 HCAPLUS
DOCUMENT NUMBER: 138:336417
TITLE: Human monoclonal antibodies and fragments specific to GEEDLP repeats for cancer diagnosis and therapy
INVENTOR(S): Takeuchi, Toshihiko; Dubois-Stringfellow, Nathalie; Murphy, John E.; Rinkenberger, Julie
PATENT ASSIGNEE(S): Bayer Corporation, USA; Bayer Pharmaceuticals Corporation
SOURCE: PCT Int. Appl., 61 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003033674	A2	20030424	WO 2002-US33470	20021018
WO 2003033674	A3	20030821		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG,

US, UZ, VC, VN, YU, ZA, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

US 2003077277	A1	20030424	US 2002-273541	20021018
EP 1438339	A2	20040721	EP 2002-786449	20021018

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, SK

PRIORITY APPLN. INFO.:	US 2001-343657P	P 20011018
	US 2002-377716P	P 20020502
	WO 2002-US33470	W 20021018

AB The invention is composed of monoclonal human MN antibodies or MN antibody fragments that target the GEEDLP repeat within the proteoglycan domain. The proteoglycan domain of the MN cell surface protein contains four of these identical GEEDLP repeats. Binding to the desired epitope is verified by competition ELISA, where ELISA signal can be attenuated by co-incubation with a peptide containing this repeat (PGEEDLPGEEDLP). This inhibition of binding can also be verified using Biacore assays, where binding of desired antibodies to immobilized MN or proteoglycan peptides can be inhibited by the peptide repeat. In addition to binding to the peptide repeat, human anti-MN antibodies can inhibit the cell adhesion of CGL-1 cells to MN coated plastic plates. Human anti-MN antibodies have been used to diagnose and quantify MN expression in cancer cells and tumors using FACS and immunohistochem. methods. An example is also provided where a human anti-MN IgG1 mediates tumor cell lysis though antibody-dependent cell-mediated cytotoxicity. Therefore, these antibodies will be useful for the treatment of cancers in which MN is upregulated or can be useful for the diagnosis of cancers in which MN is upregulated.

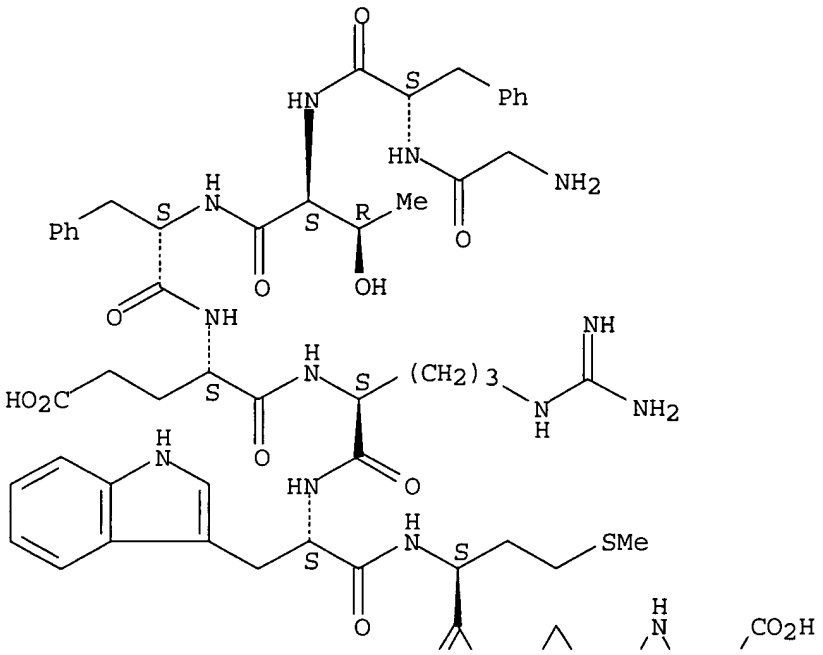
IT **517857-72-6P**
RL: BPN (Biosynthetic preparation); BSU (Biological study, unclassified); DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(human monoclonal antibodies and fragments specific to GEEDLP repeats for cancer diagnosis and therapy)

RN 517857-72-6 HCAPLUS

CN L-Alanine, glycyl-L-phenylalanyl-L-threonyl-L-phenylalanyl-L-α-glutamyl-L-arginyl-L-tryptophyl-L-methionylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



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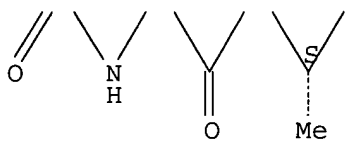


L14 ANSWER 9 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:888494 HCAPLUS
DOCUMENT NUMBER: 137:381503
TITLE: Compositions and methods for modulating Dkk-mediated protein interactions and their diagnostic and therapeutic uses
INVENTOR(S): Allen, Kristina; Anisowicz, Anthony; Bhat, Bheem M.; Damagnez, Veronique; Robinson, John Allen; Yaworsky, Paul J.
PATENT ASSIGNEE(S): Genome Therapeutics Corporation, USA; Wyeth, John and Brother Ltd.
SOURCE: PCT Int. Appl., 376 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 4
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

WO 2002092015 A2 20021121 WO 2002-US15982 20020517
WO 2002092015 A3 20031023
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
EP 1395285 A2 20040310 EP 2002-744162 20020517
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
BR 2002009836 A 20041207 BR 2002-9836 20020517
US 2004038860 A1 20040226 US 2002-182936 20020802
PRIORITY APPLN. INFO.: US 2001-291311P P 20010517
US 2002-353058P P 20020201
US 2002-361293P P 20020304
WO 2002-US15982 W 20020517

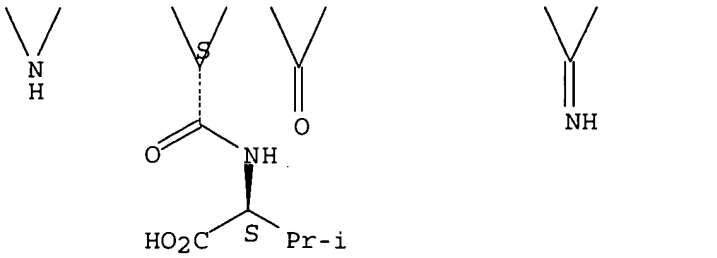
AB The present invention provides reagents, compds., compns., and methods relating to interactions of the extracellular domain of LRP5/ZMax1, HBM (a variant of LRP5), and/or LRP6 with Dkk, including Dkk-1. The various nucleic acids, polypeptides, antibodies, assay methods, diagnostic methods, and methods of treatment of the present invention are related to and impact on Dkk, LRP5, LRP6, HBM, and Wnt signaling. The invention claims sequences for peptide aptamers which bind to LRP5 or Dkk-1 and sequences for Dkk-1 peptides which are recognized by antibodies. HBM is a Gly171Val polymorphism in LDL receptor-related protein LRP5/Zmax, which has been identified as conferring a high bone mass phenotype in a population of related humans. The protein dickkopf-1 (Dkk-1) is required for head formation in early development and murine limb morphogenesis and is reported to be an antagonist of the Wnt signaling pathway. Dkk-1 protein interacts with the ligand-binding domain of LRP5. Dkk-1 also binds to LRP6, but the EGF repeat domains of LRP6 rather than the ligand-binding domain are required for interaction. Dkk-1 is able to repress LRP5-mediated Wnt signaling but not HBM-mediated Wnt signaling and Dkk-1 also inhibits LRP6 activity. LRP5, LRP6, HBM, Dkk and Wnt are implicated in bone and lipid cellular signaling. Thus, the present invention provides reagents and methods for modulating lipid levels and/or bone mass and is useful in the treatment and diagnosis of abnormal lipid levels and bone mass disorders, such as osteoporosis. Examples of the invention include a yeast two-hybrid screen for Dkk-1 interacting proteins, generation of LRP5 polymorphism-specific antibodies and Dkk-1 specific antibodies, effects of exogenous Dkk-1 on Wnt-mediated signaling in the Xenopus embryo assay, and effects of recombinant Dkk and Wnt3a/1 on TCF-luciferase reporter gene expression in human cell lines with endogenous LRP5/6.

IT **476153-38-5**
RL: DGN (Diagnostic use); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(LRP5 ligand-binding domain-interacting peptide aptamer; compns. and methods for modulating Dkk-mediated protein interactions and their diagnostic and therapeutic uses)

RN 476153-38-5 HCAPLUS

CN L-Valine, glycyl-L-tryptophyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl-L-arginyl-L-cysteinylglycyl-L-alanyl-L-leucyl-L-tryptophyl-L-tryptophyl-L-arginyl-L-arginyl- (9CI) (CA INDEX NAME)

PAGE 2-B



L14 ANSWER 10 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:382316 HCAPLUS
DOCUMENT NUMBER: 137:16533
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human lung
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 634 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001086003	A2	20011115	WO 2001-XF665	20010130
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
GB 2360284	A1	20010919	GB 2000-24263	20001004
GB 2360284	B2	20020227		
GB 2361238	A1	20011017	GB 2001-15281	20001004
GB 2361238	B2	20020306		
WO 2001086003	A2	20011115	WO 2001-US665	20010130
WO 2001086003	A3	20030522		
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GB 2396351	A1	20040623	GB 2004-6165	20010130
GB 2396351	B2	20040825		

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GB 2396352	B2	20040825			
GB 2397376	A1	20040721	GB 2004-8566		20010130
GB 2397376	B2	20041201			
US 2002102252	A1	20020801	US 2001-827998		20010406
US 6656700	B2	20031202			
US 2004063134	A1	20040401	US 2003-675685		20030930
PRIORITY APPLN. INFO.:			US 2000-180312P	P	20000204
			US 2000-207456P	P	20000526
			US 2000-608408	A	20000630
			US 2000-632366	A	20000803
			US 2000-234687P	P	20000921
			US 2000-236359P	P	20000927
			GB 2000-24263	A	20001004
			WO 2001-US665	W	20010130
			GB 2002-16928	A3	20010130
			GB 2002-17714	A3	20010130
			GB 2002-18673	A3	20010130
			US 2001-827998	A3	20010406

AB A single exon nucleic acid microarray comprising 12,614 single exon nucleic acid probes for measuring gene expression in a sample derived from human lung cells is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human lung. Also described are 12,386 single exon nucleic acid probes and 12,011 proteins expressed in the lung and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of nine records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

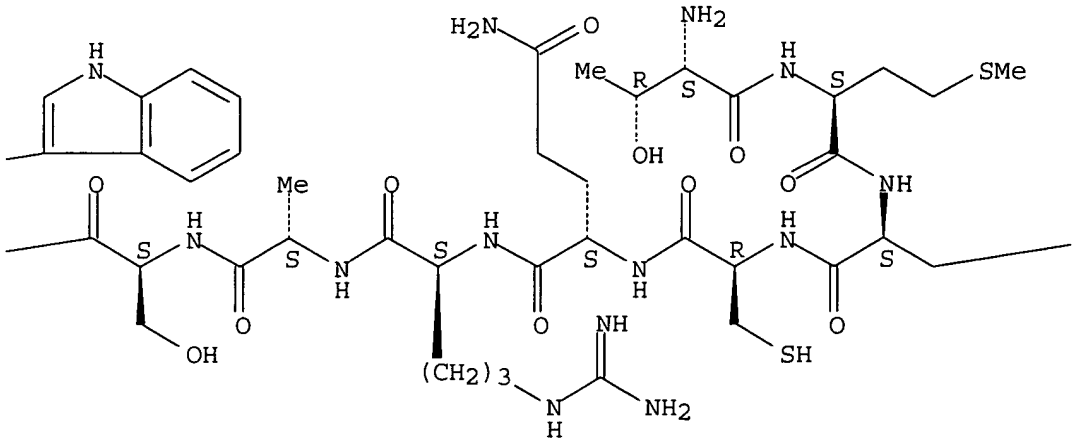
IT **400620-38-4**
RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human lung)

RN 400620-38-4 HCAPLUS

CN L-Arginine, L-threonyl-L-methionyl-L-tryptophyl-L-cysteinyl-L-glutaminyl-L-arginyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-threonyl-L-alanyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-tryptophyl-L-methionylglycyl-L-glutaminyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-C



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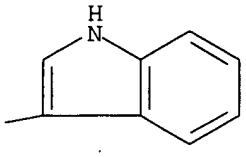

L14 ANSWER 11 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:348599 HCAPLUS
DOCUMENT NUMBER: 137:28982
TITLE: Human genome derived single exon nucleic acid probes useful for gene expression analysis
INVENTOR(S): Penn, Sharron Gaynor; Rank, David Russell; Chen, Wensheng; Hanzel, David Kagen
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 97 pp., Cont.-in-part of U.S. Ser. No. 774,203.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002048763	A1	20020425	US 2001-864761	20010523
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PRIORITY APPLN. INFO.:

US 2000-180312P	P	20000204
US 2000-207456P	P	20000526
US 2000-608408	A2	20000630
US 2000-632366	A2	20000803
US 2000-234687P	P	20000921
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GB 2002-17714	A3	20010130
GB 2002-18673	A3	20010130
US 2001-266860P	P	20010205
US 2001-827998	A3	20010406

AB Methods and apparatus for predicting, confirming and displaying functional regions from genomic sequence data are used to identify 16,834 unique human genome-derived single exon probes useful for gene expression anal., particularly gene expression anal. by microarray. Also presented are genome-derived single exon microarrays that include such probes, peptides encoded by the exons, and antibodies thereto. The human genome-derived single-exon probes are known to be expressed in one or more human tissues or cell types, particularly human brain, heart, liver, fetal liver,placenta, lung, bone marrow, BT474 and other human mammary epithelial cells, HeLa and other human cervical epithelial cells, and HBL 100 and other human mammary epithelial cells. The invention provides a method of financing, selling and/or licensing genome-derived single-exon microarrays to customer desiring to measure gene expression, comprising: making available for computerized query or subscription service a database having a record corresponding to each genome-derived single exon microarray available for sale and/or license. [This abstract record is one of ten records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

IT **400620-38-4**
RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)
(amino acid sequence; human genome derived single exon nucleic acid probes useful for gene expression anal.)

RN 400620-38-4 HCAPLUS

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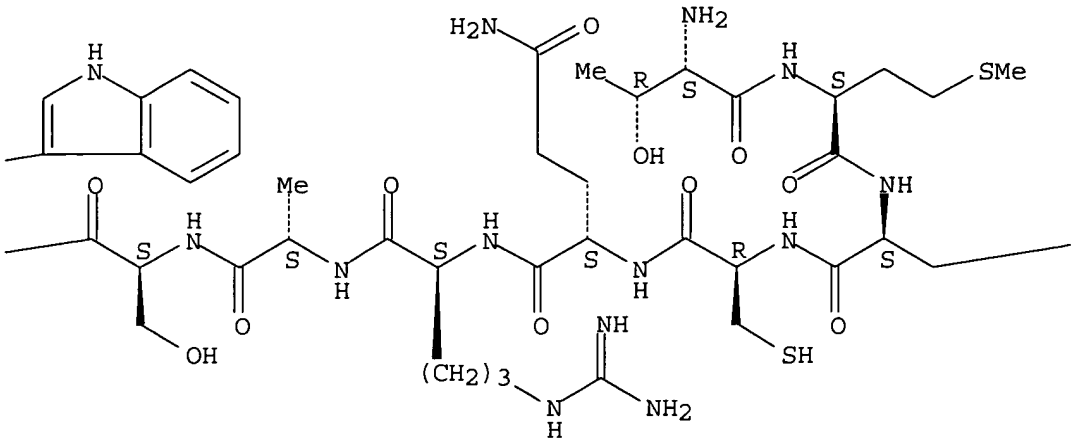

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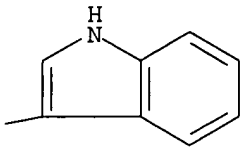

L14 ANSWER 12 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:173785 HCAPLUS
DOCUMENT NUMBER: 136:351355
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human adult liver
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 658 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001057273	A2	20010809	WO 2001-XF664	20010130

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GB 2397376	A1	20040721	GB 2004-8566	20010130
GB 2397376	B2	20041201		
US 2002102252	A1	20020801	US 2001-827998	20010406
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PRIORITY APPLN. INFO.:

US 2000-180312P	P	20000204
US 2000-207456P	P	20000526
US 2000-608408	A	20000630
US 2000-632366	A	20000803
US 2000-234687P	P	20000921
US 2000-236359P	P	20000927
GB 2000-24263	A	20001004
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GB 2002-16928	A3	20010130
GB 2002-17714	A3	20010130
GB 2002-18673	A3	20010130
US 2001-827998	A3	20010406

AB A single exon nucleic acid microarray comprising 13,109 single exon nucleic acid probes for measuring gene expression in a sample derived from human adult liver is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human adult liver cells. Also described are 12,886 single exon nucleic acid probes and 12,583 proteins expressed in the adult liver and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene.

IT **400620-38-4**
RL: ANT (Analyte); BSU (Biological study, unclassified); BUU (Biological

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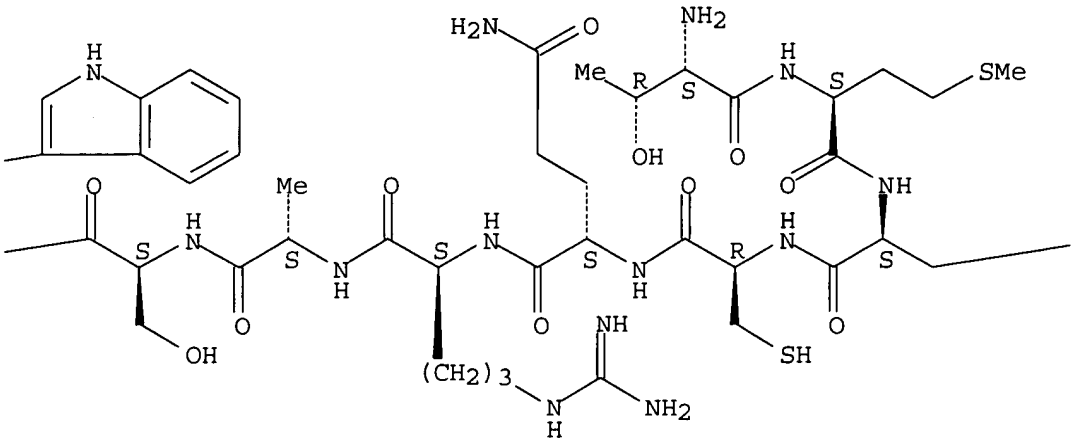

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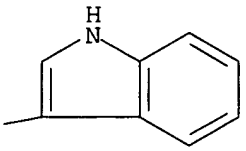

L14 ANSWER 13 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:110620 HCAPLUS
DOCUMENT NUMBER: 136:195269
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human placenta
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 654 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001057272	A2	20010809	WO 2001-XD663	20010130

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required to fully index the document and publication system constraints.].

IT **400620-38-4**
RL: ANT (Analyte); BSU (Biological study, unclassified); PRP (Properties); ANST (Analytical study); BIOL (Biological study)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human placenta)

RN 400620-38-4 HCAPLUS

CN L-Arginine, L-threonyl-L-methionyl-L-tryptophyl-L-cysteinyl-L-glutaminyl-L-arginyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-threonyl-L-alanyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-tryptophyl-L-methionylglycyl-L-glutaminyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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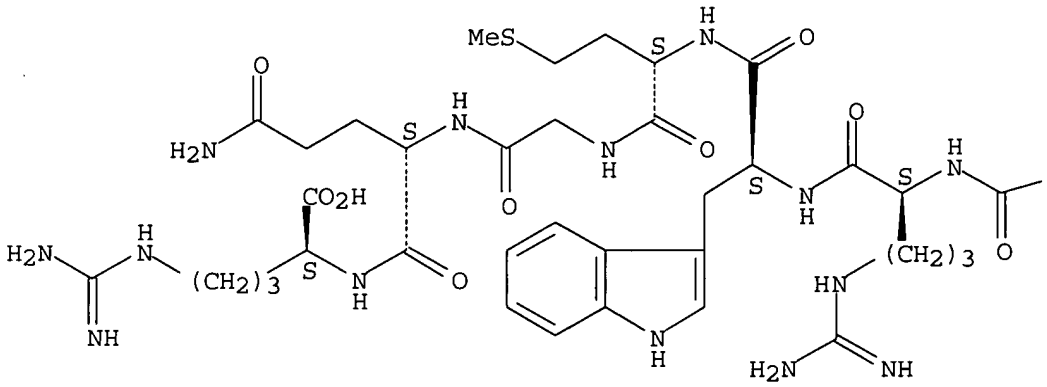

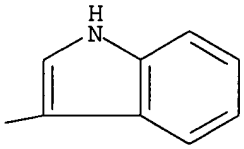

L14 ANSWER 14 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:110610 HCAPLUS
DOCUMENT NUMBER: 136:351347
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human HeLa cells or other human cervical epithelial cells
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 487 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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			GB 2002-17714	A3	20010130
			GB 2002-18673	A3	20010130
			US 2001-827998	A3	20010406

AB A single exon nucleic acid microarray comprising 9290 single exon nucleic acid probes for measuring gene expression in a sample derived from human HeLa cells or other human cervical epithelial cells is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human HeLa cells or other human cervical epithelial cell lines. Also described are 9102 single exon nucleic acid probes and 8549 proteins expressed in the cervical epithelial cells and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of six records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

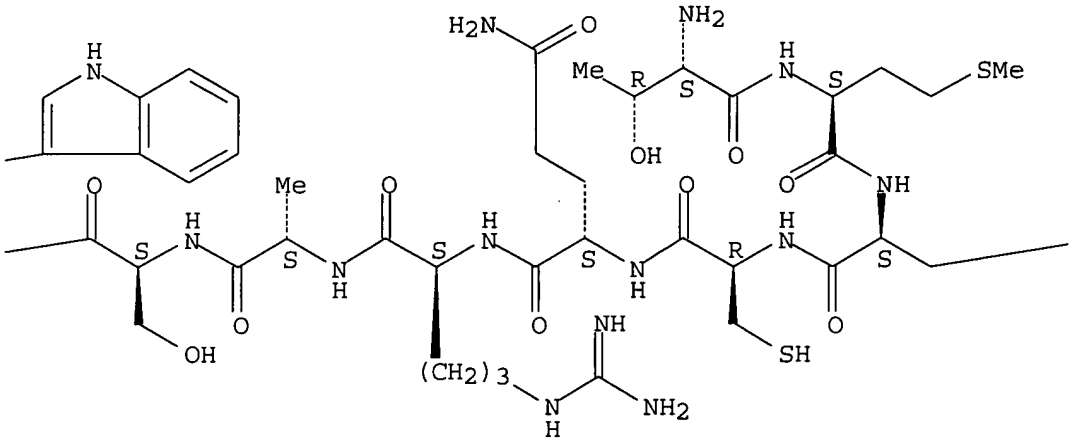
IT **400620-38-4**
RL: ANT (Analyte); BUU (Biological use, unclassified); PRP (Properties); ANST (Analytical study); BIOL (Biological study); USES (Uses)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human HeLa cells or other human cervical epithelial cells)

RN 400620-38-4 HCAPLUS

CN L-Arginine, L-threonyl-L-methionyl-L-tryptophyl-L-cysteinyl-L-glutaminyl-L-arginyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-threonyl-L-alanyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-tryptophyl-L-methionylglycyl-L-glutaminyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-C



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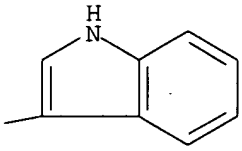

L14 ANSWER 15 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:110604 HCAPLUS
DOCUMENT NUMBER: 136:178933
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human fetal liver
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 639 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001057277	A2	20010809	WO 2001-XD669	20010130

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BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

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WO 2001057277	A3	20030213		

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GB 2396352	B2	20040825		
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GB 2397376	B2	20041201		
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US 6656700	B2	20031202		
US 2004063134	A1	20040401	US 2003-675685	20030930

PRIORITY APPLN. INFO.:

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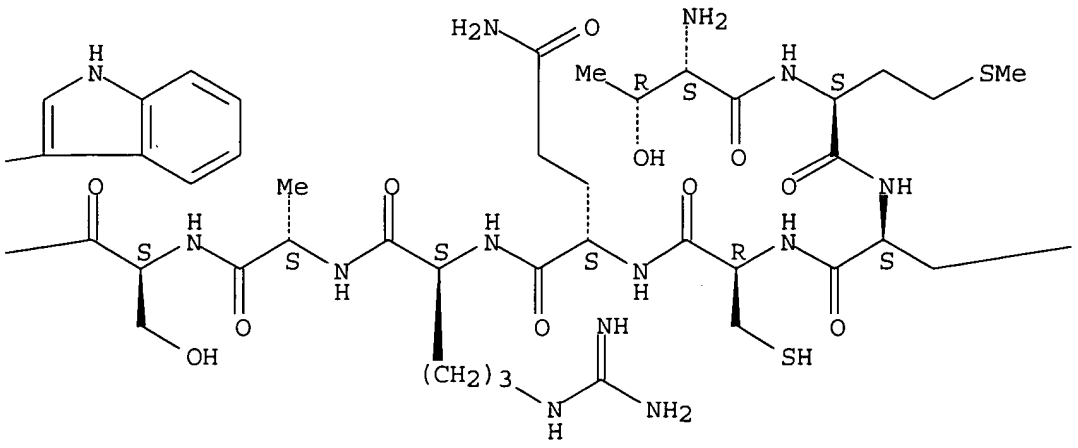
GB 2002-16928	A3	20010130
GB 2002-17714	A3	20010130
GB 2002-18673	A3	20010130
US 2001-827998	A3	20010406

AB A single exon nucleic acid microarray comprising 12,673 single exon nucleic acid probes for measuring gene expression in a sample derived from human fetal liver cells is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human fetal liver cells. Also described are 12,456 single exon nucleic acid probes and 12,027 proteins expressed in the fetal liver and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of nine records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

IT **400620-38-4**

PAGE 1-C



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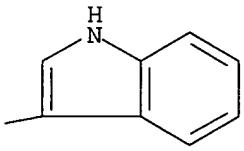

L14 ANSWER 16 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:110588 HCAPLUS
DOCUMENT NUMBER: 136:305084
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human brain
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 650 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001057275	A2	20010809	WO 2001-XD667	20010130

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GW, ML, MR, NE, SN, TD, TG

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	GB 2002-16928	A3 20010130
	GB 2002-17714	A3 20010130
	GB 2002-18673	A3 20010130
	US 2001-827998	A3 20010406

AB A single exon nucleic acid microarray comprising 12,821 single exon nucleic acid probes for measuring gene expression in a sample derived from human brain cells is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human brain. Also described are 12,613 single exon nucleic acid probes and 12,377 proteins expressed in the brain and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of nine records for

this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

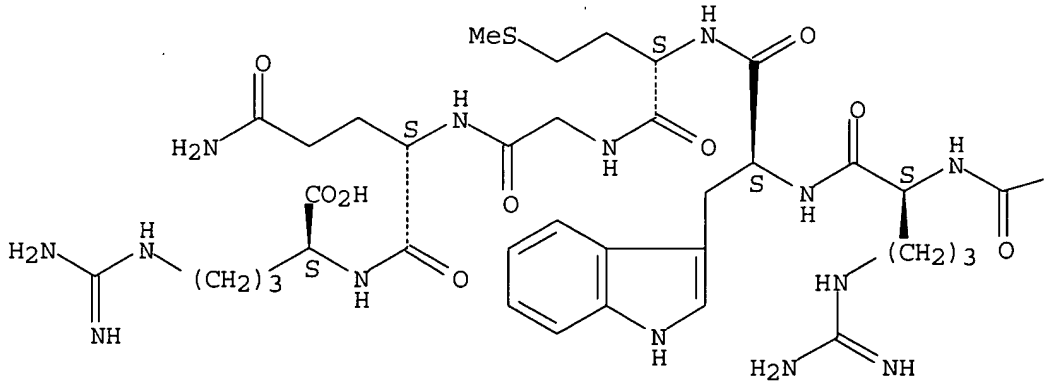
IT **400620-38-4**
RL: ANT (Analyte); BSU (Biological study, unclassified); PRP (Properties); ANST (Analytical study); BIOL (Biological study)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human brain)

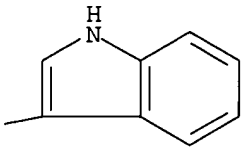
RN 400620-38-4 HCAPLUS

CN L-Arginine, L-threonyl-L-methionyl-L-tryptophyl-L-cysteinyl-L-glutaminyl-L-arginyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-threonyl-L-alanyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-tryptophyl-L-methionylglycyl-L-glutaminyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A





L14 ANSWER 17 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:110579 HCAPLUS
DOCUMENT NUMBER: 136:211829
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human bone marrow
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 657 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

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WO 2001057276	A2	20010809	WO 2001-XD668	20010130
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GB 2360284	B2	20020227		
GB 2361238	A1	20011017	GB 2001-15281	20001004
GB 2361238	B2	20020306		
WO 2001057276	A2	20010809	WO 2001-US668	20010130
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			GB 2002-17714	A3	20010130
			GB 2002-18673	A3	20010130
			US 2001-827998	A3	20010406

AB A single exon nucleic acid microarray comprising 13,114 single exon nucleic acid probes for measuring gene expression in a sample derived from human bone marrow is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human bone marrow. Also described are 12,898 single exon nucleic acid probes and 12,616 proteins expressed in the bone marrow and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of nine records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

IT **400620-38-4**
RL: ANT (Analyte); BSU (Biological study, unclassified); PRP (Properties); ANST (Analytical study); BIOL (Biological study)
(amino acid sequence; human genome-derived single exon nucleic acid probes useful for anal. of gene expression in human bone marrow)

RN 400620-38-4 HCAPLUS

CN L-Arginine, L-threonyl-L-methionyl-L-tryptophyl-L-cysteinyl-L-glutaminyl-L-arginyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-threonyl-L-alanyl-L-alanyl-L-seryl-L-tryptophyl-L-arginyl-L-tryptophyl-L-methionylglycyl-L-glutaminyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

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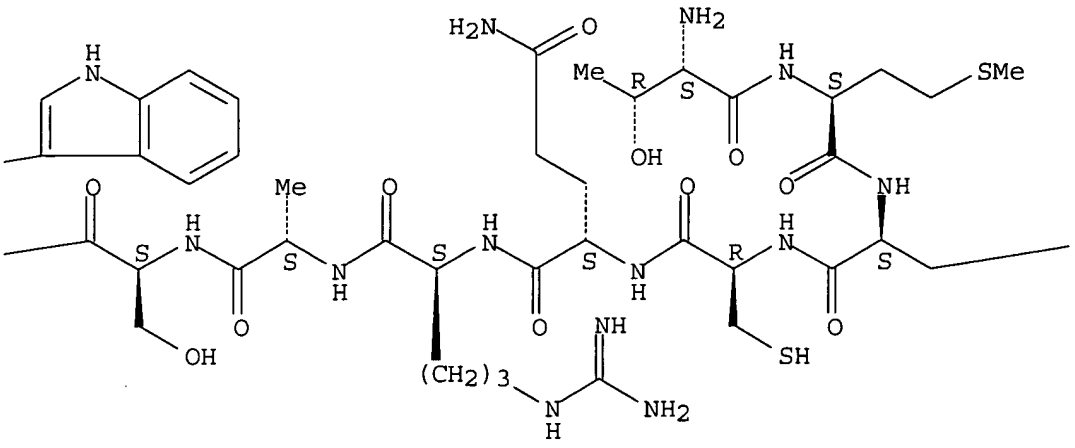

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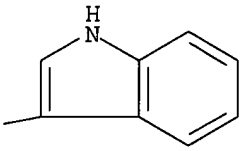

L14 ANSWER 18 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2002:64569 HCAPLUS
DOCUMENT NUMBER: 136:195264
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human heart
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 529 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001057274	A2	20010809	WO 2001-XD666	20010130

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GB 2361238	B2	20020306		
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PRIORITY APPLN. INFO.:

US 2000-180312P	P	20000204
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GB 2002-16928	A3	20010130
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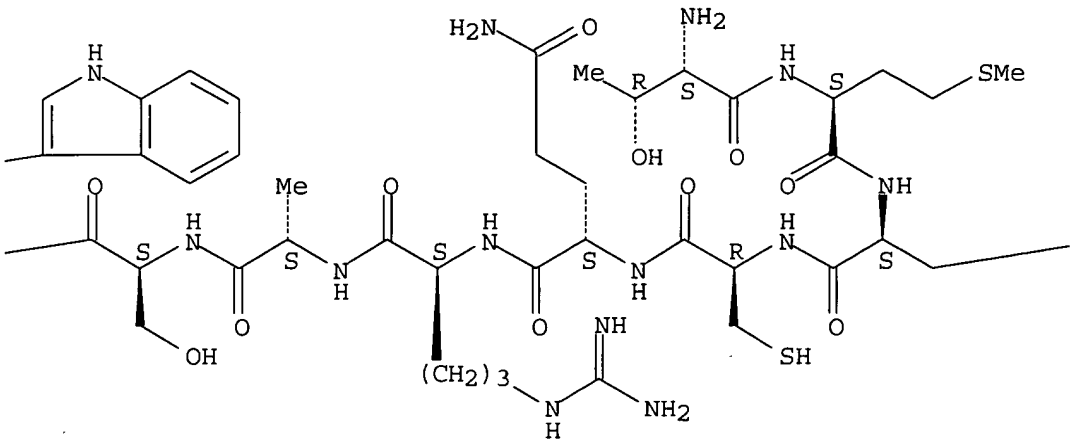
AB A single exon nucleic acid microarray comprising 9980 single exon nucleic acid probes for measuring gene expression in a sample derived from human heart is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human heart cells. Also described are 9791 single exon nucleic acid probes and 9347 proteins expressed in the heart and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of six records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

IT **400620-38-4**

PAGE 1-C



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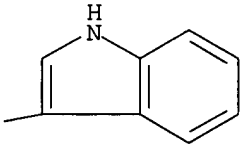

L14 ANSWER 19 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2001:934560 HCAPLUS
Correction of: 2001:582102
DOCUMENT NUMBER: 136:195263
Correction of: 135:163340
TITLE: Human genome-derived single exon nucleic acid probes useful for analysis of gene expression in human breast and BT 474 cells
INVENTOR(S): Penn, Sharron G.; Hanzel, David K.; Chen, Wensheng; Rank, David R.
PATENT ASSIGNEE(S): Molecular Dynamics, Inc., USA
SOURCE: PCT Int. Appl., 327 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 90
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

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			US 2001-327898P	P	20011009
			US 2001-335941P	P	20011024

AB A single exon nucleic acid microarray comprising 5205 single exon nucleic acid probes for measuring gene expression in a sample derived from human BT 474 cells is described. These unique exons are within longer probe sequences; sequencing confirms the exact chemical structure of each probe. Some amplicons have more than one exon, and some exons are contained in more than one amplicon. Expression, homol., and functional information are provided for the genome-derived single exon probes that are expressed significantly in human BT 474 cells. Also described are 5112 single exon nucleic acid probes and 5121 proteins expressed in the BT 474 cells and their use in methods for detecting gene expression. The genome-derived single exon nucleic acids comprise a novel type of nucleic acid microarray for verifying gene expression. In addition, methods are provided for identifying exons in a eukaryotic genome, and for assigning exons to a single gene. [This abstract record is one of three records for this document necessitated by the large number of index entries required to fully index the document and publication system constraints.].

IT **400620-38-4**
RL: BSU (Biological study, unclassified); PRP (Properties); BIOL (Biological study)

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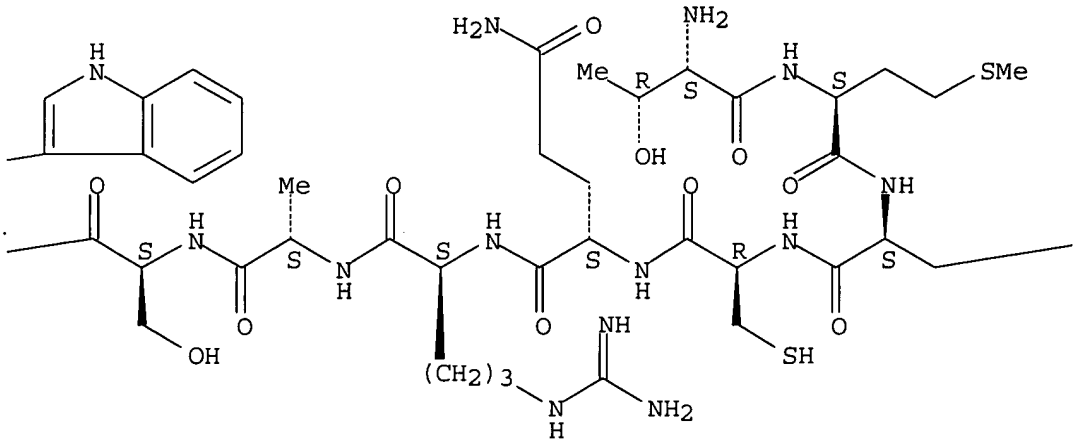

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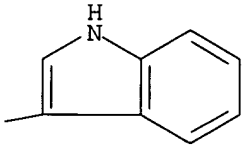

L14 ANSWER 20 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2001:816705 HCAPLUS
DOCUMENT NUMBER: 135:366701
TITLE: Fc-domain-modified peptides as therapeutic agents
INVENTOR(S): Feige, Ulrich; Liu, Chuan-Fa; Cheetham, Janet C.; Boone, Thomas Charles; Gudas, Jean Marie
PATENT ASSIGNEE(S): Amgen Inc., USA
SOURCE: PCT Int. Appl., 176 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001083525	A2	20011108	WO 2001-US14310	20010502
WO 2001083525	A3	20020718		

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AB The present invention concerns fusion of Fc domains with biol. active peptides and a process for preparing pharmaceutical agents using biol. active peptides. In this invention, pharmacol. active compds. are prepared by a process comprising: a) selecting at least one peptide that modulates the activity of a protein of interest; and b) preparing a pharmacol. agent comprising an Fc domain covalently linked to at least one amino acid of the selected peptide. Linkage to the vehicle increases the half-life of the peptide, which otherwise would be quickly degraded in vivo. The preferred vehicle is an Fc domain. The peptide can be selected, for example, by phage display, E.coli display, ribosome display, RNA-peptide screening, yeast-based screening, chemical-peptide screening, rational design, or protein structural anal.

IT **268228-13-3**
RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(Fc-domain-modified peptides as therapeutic agents)

RN 268228-13-3 HCAPLUS

CN L-Serine, L-seryl-L-cysteinyl-L-tyrosyl-L-α-glutamyl-L-tryptophylglycyl-L-lysyl-L-leucyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

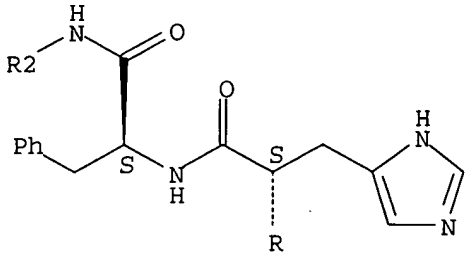
Absolute stereochemistry.

PAGE 2-A

L14 ANSWER 21 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2000:780431 HCAPLUS
DOCUMENT NUMBER: 134:68132
TITLE: Melanoma-targeting properties of 99mtechnetium-labeled cyclic α-melanocyte-stimulating hormone peptide analogues
AUTHOR(S): Chen, JianQing; Cheng, Zhen; Hoffman, Timothy J.; Jurisson, Silvia S.; Quinn, Thomas P.
CORPORATE SOURCE: Department of Biochemistry, University of Missouri-Columbia, Columbia, MO, 65211, USA
SOURCE: Cancer Research (2000), 60(20), 5649-5658
CODEN: CNREA8; ISSN: 0008-5472
PUBLISHER: American Association for Cancer Research
DOCUMENT TYPE: Journal
LANGUAGE: English

AB Preliminary reports have demonstrated that 99mtechnetium (Tc)-labeled cyclic [Cys3,4,10, D-Phe7]α-MSH3-13 (CCMSH) exhibits high tumor uptake and retention values in a murine melanoma mouse model. In this report, the tumor targeting mechanism of 99mTc-CCMSH was studied and compared with four other radiolabeled α-MSH (α-MSH) peptide analogs: 125I-(Tyr2)-[Nle4, D-Phe7]α-MSH [125I-(Tyr2)-NDP]; 99mTc-CGCG-NDP; 99mTc-Gly11-CCMSH; and 99mTc-Nle11-CCMSH. In vitro receptor binding, internalization, and cellular retention of radiolabeled α-MSH analogs in B16/F1 murine cell line demonstrated that >70% of the receptor-bound radiolabeled analogs were internalized together with the receptor. Ninety % of the internalized 125I-(Tyr2)-NDP, whereas only 36% of internalized 99mTc-CCMSH, was released from the cells into the medium during a 4-h incubation at 37°C. Two mouse models, C57 mice and severe combined immunodeficient (Scid) mice, inoculated s.c. with B16/F1 murine and TXM-13 human melanoma cells were used for the in vivo studies. Tumor uptake values of 11.32 and 2.39 [% injected dose (ID)/g] for 99mTc-CCMSH at 4 h after injection, resulted in an uptake ratio of tumor:blood of 39.0 and 11.5 in murine melanoma-C57 and human melanoma-Scid mouse models, resp. Two strategies for decreasing the nonspecific kidney uptake of 99mTc-CCMSH, substitution of Lys11 in CCMSH with Gly11 or Nle11, and lysine coinjection, were evaluated. The biodistribution data for the modified peptides showed that Lys11 replacement dramatically decreased the kidney uptake, whereas the tumor uptakes of 99mTc-Nle11- and 99mTc-Gly11-CCMSH were significantly lower than that of 99mTc-CCMSH. Lysine coinjection significantly decreased the kidney uptake (e.g., from 14.6% ID/g to 4.5% ID/g at 4 h after injection in murine melanoma-C57 mice) without significantly changing the value of tumor uptake of 99mTc-CCMSH. In conclusion, the compact cyclic structure of 99mTc-CCMSH, its resistance to degradation, and its enhanced intracellular retention are the major contributing factors to the superior in vivo tumor targeting properties of 99mTc-CCMSH. Lys11 residue in 99mTc-CCMSH is critical to the tumor targeting in vivo, and lysine coinjection rather than lysine replacement can significantly decrease the nonspecific renal radioactivity accumulation without impeding the high melanoma-targeting properties of 99mTc-CCMSH. The metal-cyclized CCMSH mol. displays excellent potential for the development of melanoma-specific diagnostic and therapeutic agents.

PAGE 3-A



REFERENCE COUNT: 40 THERE ARE 40 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 22 OF 22 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2000:291095 HCAPLUS
DOCUMENT NUMBER: 132:329919
TITLE: Modified peptides containing an antibody Fc domain as therapeutic agents
INVENTOR(S): Feige, Ulrich; Liu, Chuan-fa; Cheetham, Janet; Boone, Thomas Charles
PATENT ASSIGNEE(S): Amgen Inc., USA
SOURCE: PCT Int. Appl., 608 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2000024782	A2	20000504	WO 1999-US25044	19991025
WO 2000024782	A3	20020606		
W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
US 6660843	B1	20031209	US 1999-428082	19991022
CA 2347131	AA	20000504	CA 1999-2347131	19991025
EP 1144454	A2	20011017	EP 1999-971003	19991025
EP 1144454	A3	20020911		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
BR 9914708	A	20020716	BR 1999-14708	19991025
JP 2003512011	T2	20030402	JP 2000-578351	19991025
AU 767725	B2	20031120	AU 2000-12322	19991025
NZ 510888	A	20040130	NZ 1999-510888	19991025
ZA 2001002753	A	20020611	ZA 2001-2753	20010404
NO 2001001963	A	20010621	NO 2001-1963	20010420
BG 105461	A	20030430	BG 2001-105461	20010424
US 2004044188	A1	20040304	US 2003-609217	20030627
US 2004053845	A1	20040318	US 2003-632388	20030731

US 2004071712	A1	20040415	US 2003-645761	20030818
US 2004057953	A1	20040325	US 2003-651723	20030829
US 2004087778	A1	20040506	US 2003-653048	20030829
US 2004077022	A1	20040422	US 2003-666696	20030919
PRIORITY APPLN. INFO.:			US 1998-105371P	P 19981023
			US 1999-428082	A 19991022
			WO 1999-US25044	W 19991025
			US 2000-563286	A1 20000503

AB The present invention concerns fusion of Fc domains with biol. active peptides and a process for preparing pharmaceutical agents using biol. active peptides. In this invention, pharmacol. active compds. are prepared by a process comprising: (a) selecting at least one peptide that modulates the activity of a protein of interest; and (b) preparing a pharmacol. agent comprising an Fc domain covalently linked to at least one amino acid of the selected peptide. Linkage to the vehicle increases the half-life of the peptide, which otherwise would be quickly degraded in vivo. The preferred vehicle is an Fc domain. The peptide is preferably selected by phage display, Escherichia coli coli display, ribosome display, RNA-peptide screening, or chemical-peptide screening.

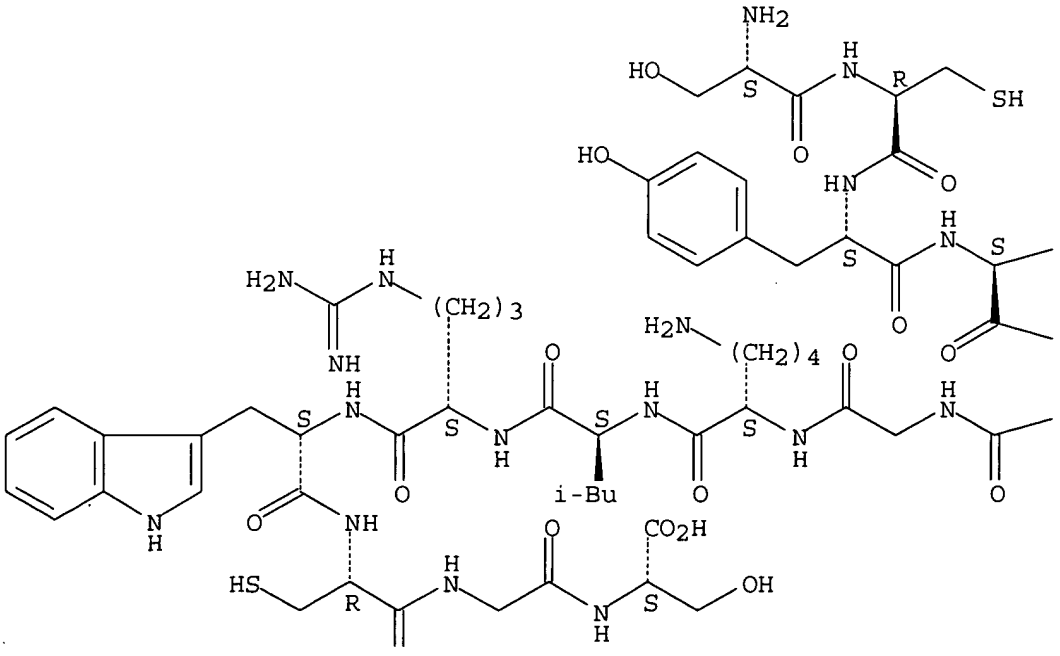
IT **268228-13-3D**, fusion protein with IgG1 Fc domain
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(calmodulin antagonist; modified peptides containing an antibody Fc domain as therapeutic agents)

RN 268228-13-3 HCAPLUS

CN L-Serine, L-seryl-L-cysteinyl-L-tyrosyl-L-α-glutamyl-L-tryptophylglycyl-L-lysyl-L-leucyl-L-arginyl-L-tryptophyl-L-cysteinylglycyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

PAGE 1-A



=> d ibib abs hitstr l29 1-3

L29 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 2004:995718 HCAPLUS
DOCUMENT NUMBER: 141:416010
TITLE: Erythropoietin conjugate compounds with extended half-lives
INVENTOR(S): **Heavner, George**
PATENT ASSIGNEE(S): USA
SOURCE: U.S. Pat. Appl. Publ., 11 pp.
CODEN: USXXCO
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

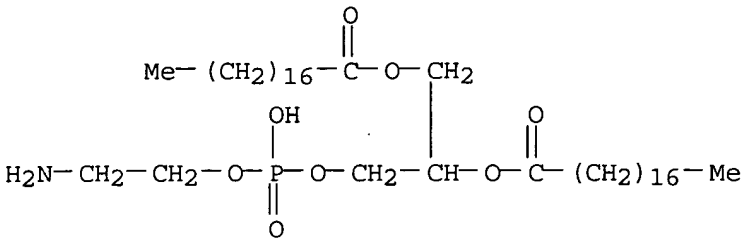
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2004229318	A1	20041118	US 2003-439870	20030517
WO 2004106373	A1	20041209	WO 2003-US15750	20030520

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR

PRIORITY APPLN. INFO.: US 2003-439870 A 20030517

AB The invention provides biol. active erythropoietin (EPO) conjugate compns. wherein EPO is covalently conjugated to a non-antigenic hydrophilic polymer covalently linked to an organic mol. that increases the circulating serum half-life of the composition The invention thus relates to EPO derivs. described by the formula EPO-(X-Y) N where EPO is erythropoietin or its pharmaceutical acceptable derivs. having biol. properties of causing **bone** marrow cells to increase production of reticulocytes and red blood cells, X is PEG or other water soluble polymers, Y is an organic mol. that increases the circulating half-life of the construct more than the PEG alone and N is an integer from 1 to 15. Other mols. may be included between EPO and X and between X and Y to provide the proper functionality for coupling or valency. For example, erythropoietin was conjugated to DSPE-PEG through the alpha amino group of amino acid 1 of erythropoietin, and was able to prolong the serum half-life of erythropoietin in mice shown by the high hematocrit and Hb levels.

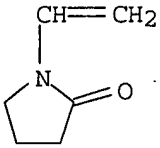
IT **76-05-1**, Trifluoroacetic acid, uses **7087-68-5**, Diisopropylethylamine
RL: NUU (Other use, unclassified); USES (Uses)
(erythropoietin conjugates with polymers and orgs. for extended serum half-lives)
RN 76-05-1 HCAPLUS
CN Acetic acid, trifluoro- (8CI, 9CI) (CA INDEX NAME)



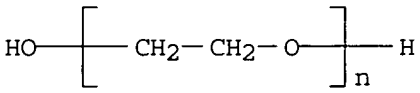
RN 9003-39-8 HCAPLUS
CN 2-Pyrrolidinone, 1-ethenyl-, homopolymer (9CI) (CA INDEX NAME)

CM 1

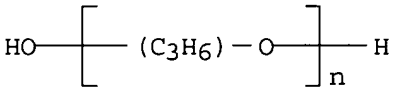
CRN 88-12-0
CMF C6 H9 N O



RN 25322-68-3 HCAPLUS
CN Poly(oxy-1,2-ethanediyl), α-hydro-ω-hydroxy- (9CI) (CA INDEX NAME)



RN 25322-69-4 HCAPLUS
CN Poly[oxy(methyl-1,2-ethanediyl)], α-hydro-ω-hydroxy- (9CI) (CA INDEX NAME)



L29 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2005 ACS on STN
ACCESSION NUMBER: 1986:573063 HCAPLUS
DOCUMENT NUMBER: 105:173063
TITLE: Bursopoietin
INVENTOR(S): Audhya, Tapan; Kroon, Daniel J.; **Heavner, George**; Goldstein, Gideon
PATENT ASSIGNEE(S): Ortho Pharmaceutical Corp., USA
SOURCE: U.S., 6 pp.
CODEN: USXXAM
DOCUMENT TYPE: Patent
LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2
PATENT INFORMATION:

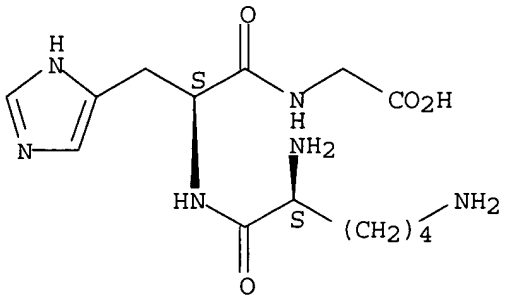
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 4584284	A	19860422	US 1985-681971	19850131
AU 8652737	A1	19860807	AU 1986-52737	19860124
AU 593231	B2	19900208		
ES 551399	A1	19870516	ES 1986-551399	19860129
IL 77736	A1	19900429	IL 1986-77736	19860129
DK 8600476	A	19860801	DK 1986-476	19860130
EP 190048	A2	19860806	EP 1986-300631	19860130
EP 190048	A3	19880803		
EP 190048	B1	19920122		
R: AT, BE, CH, DE, FR, GB, IT, LI, LU, NL, SE				
JP 61218598	A2	19860929	JP 1986-16965	19860130
ZA 8600702	A	19870826	ZA 1986-702	19860130
AT 71953	E	19920215	AT 1986-300631	19860130
CA 1276398	A1	19901113	CA 1986-500890	19860131
US 4783442	A	19881108	US 1986-855011	19860421
US 4866121	A	19890912	US 1988-211203	19880623

PRIORITY APPLN. INFO.:
US 1985-681971 A 19850131
EP 1986-300631 A 19860130
US 1986-855011 A3 19860421

AB A **peptide** (H-Lys-His-Gly-NH2) was prepared by solid-phase synthesis which induced differentiation of **bone** marrow cells to B-lymphocytes (bursopoietin-like activity).

IT **104768-75-4P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 104768-75-4 HCAPLUS
CN Glycine, N-(N-L-lysyl-L-histidyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT **60267-34-7P**
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of and B-lymphocyte differentiation response to)
RN 60267-34-7 HCAPLUS
CN Glycinamide, L-lysyl-L-histidyl- (9CI) (CA INDEX NAME)

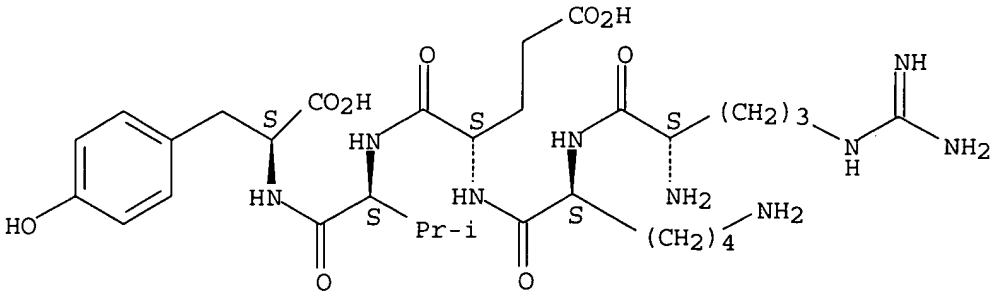
Absolute stereochemistry.

ACCESSION NUMBER: 1983:592890 HCAPLUS
DOCUMENT NUMBER: 99:192890
TITLE: Proton nuclear magnetic resonance investigation of the active site fragment of splenin, an immunoregulatory **polypeptide**
AUTHOR(S): Krishna, N. Rama; **Heavner, George A.**; Vaughn, Joseph B., Jr.
CORPORATE SOURCE: Compr. Cancer Cent., Univ. Alabama, Birmingham, AL, 35294, USA
SOURCE: Journal of the American Chemical Society (1983), 105(23), 6930-4
CODEN: JACSAT; ISSN: 0002-7863
DOCUMENT TYPE: Journal
LANGUAGE: English

AB The **peptide** fragment Arg-Lys-Glu-Val-Tyr (SP5) is the active site fragment of splenin (formerly thymopoietin III), an immunoregulatory **polypeptide** isolated from bovine spleen. An NMR investigation of the conformationaL properties of this active fragment in aqueous soln is reported. All the observed NH and CH resonances of SP5 were assigned by 1-dimensional and 2-dimensional NMR techniques. The variation of chemical shifts with pH, the individual amide hydrogen exchange rates, and the vicinal NH-CαH coupling consts. were measured. The data are compatible with the assumption of a highly motile dynamic equilibrium among different conformations, some of which are stabilized by internal H bonding involving the participation of Glu3-NH, Val4-NH, and Tyr5-NH in the **backbone** and of the guanidino NεH proton of the Arg1 side chain. These observations provide an insight into the conformational tendencies of SP5 in aqueous solns.

IT **75957-60-7**
RL: BIOL (Biological study)
(as splenin active site, NMR and conformation of)
RN 75957-60-7 HCAPLUS
CN L-Tyrosine, L-arginyl-L-lysyl-L-α-glutamyl-L-valyl- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



IT **1416-60-0**
RL: BIOL (Biological study)
(**pentapeptide** of active site of, NMR and conformation of)
RN 1416-60-0 HCAPLUS
CN Splenin (9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***